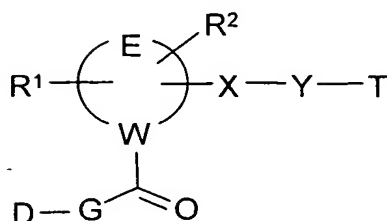


Patent Claims

1. Compounds of the formula I



in which

R^1, R^2 are each, independently of one another, H, =O, Hal, A, ethynyl, OR^3 , $\text{N}(\text{R}^3)_2$, NO_2 , CN, N_3 , COOR^3 , $\text{CON}(\text{R}^3)_2$, $-\text{[C(R}^4)_2\text{]}_n\text{-Ar}$, $-\text{[C(R}^4)_2\text{]}_n\text{-Het}$, $-\text{[C(R}^4)_2\text{]}_n\text{-cycloalkyl}$, $-\text{OCOR}^3$, $-\text{OCON}(\text{R}^3)_2$, NR^3COA or $\text{NR}^3\text{SO}_2\text{A}$,

R^1 and R^2 together are alternatively a bicyclically or spirocyclically bonded 3- to 7-membered carbocyclic or heterocyclic ring having from 0 to 3 N, O and/or S atoms,

R^3 is H, A, $\text{H-C}\equiv\text{C-CH}_2\text{-}$, $\text{CH}_3\text{-C}\equiv\text{C-CH}_2\text{-}$, $-\text{CH}_2\text{-CH(OH)-CH}_2\text{OH}$, $-\text{CH}_2\text{-CH(OH)-CH}_2\text{NH}_2$, $-\text{CH}_2\text{-CH(OH)-CH}_2\text{Het}'$, $-\text{[C(R}^4)_2\text{]}_n\text{-Ar}'$, $-\text{[C(R}^4)_2\text{]}_n\text{-Het}'$, $-\text{[C(R}^4)_2\text{]}_n\text{-cycloalkyl}$, $-\text{[C(R}^4)_2\text{]}_n\text{-COOA}$ or $-\text{[C(R}^4)_2\text{]}_n\text{N(R}^4)_2$,

R^4 is H or A,

W is N, CR^3 or an sp^2 -hybridised carbon atom,

E together with W is a 3- to 7-membered saturated carbocyclic or heterocyclic ring having from 0 to 3 N, from 0 to 2 O and/or from 0 to 2 S atoms,

which may contain a double bond,

D is a monocyclic or bicyclic, aromatic carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or polysubstituted by Hal, A, OR^3 , $\text{N(R}^3)_2$, NO_2 , CN, COOR^3 or $\text{CON(R}^3)_2$,

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- G is $-\text{C}(\text{R}^4)_2]_n-$, $-\text{C}(\text{R}^4)_2]_n\text{NR}^3-$, $-\text{C}(\text{R}^4)_2]_n\text{O}-$, $-\text{C}(\text{R}^4)_2]_n\text{S}-$ or $-\text{C}(\text{R}^4)=\text{C}(\text{R}^4)]_n-$,
- X is $-\text{C}(\text{R}^4)_2]_n\text{CONR}^3[\text{C}(\text{R}^4)_2]_n-$, $-\text{C}(\text{R}^4)_2]_n\text{NR}^3\text{CO}[\text{C}(\text{R}^4)_2]_n-$, $-\text{C}(\text{R}^4)_2]_n\text{NR}^3[\text{C}(\text{R}^4)_2]_n-$, $-\text{C}(\text{R}^4)_2]_n\text{O}[\text{C}(\text{R}^4)_2]_n-$, $-\text{C}(\text{R}^4)_2]_n\text{CO}[\text{C}(\text{R}^4)_2]_n-$ or $-\text{C}(\text{R}^4)_2]_n\text{COO}[\text{C}(\text{R}^4)_2]_n-$,
- Y is alkylene, cycloalkylene, Het-diyl or Ar-diyl,
- T is a monocyclic or bicyclic, saturated or unsaturated carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is monosubstituted or disubstituted by $=\text{O}$, $=\text{S}$, $=\text{NR}^3$, $=\text{N-CN}$, $=\text{N-NO}_2$, $=\text{NOR}^3$, $=\text{NCOR}^3$, $=\text{NCOOR}^3$ or $=\text{NOCOR}^3$ and may furthermore be monosubstituted, disubstituted or trisubstituted by R^3 , Hal, A, $-\text{C}(\text{R}^4)_2]_n\text{-Ar}$, $-\text{C}(\text{R}^4)_2]_n\text{-Het}$, $-\text{C}(\text{R}^4)_2]_n\text{-cycloalkyl}$, OR^3 , $\text{N}(\text{R}^3)_2$, NO_2 , CN , COOR^3 , $\text{CON}(\text{R}^3)_2$, NR^3COA , $\text{NR}^3\text{CON}(\text{R}^3)_2$, $\text{NR}^3\text{SO}_2\text{A}$, COR^3 , SO_2NR^3 and/or $\text{S}(\text{O})_n\text{A}$,
- A is unbranched or branched alkyl having 1-10 carbon atoms in which one or two CH_2 groups may be replaced by O or S atoms and/or by $-\text{CH}=\text{CH}-$ groups and/or in addition 1-7 H atoms may be replaced by F,
- Ar is phenyl, naphthyl or biphenyl, each of which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR^3 , $\text{N}(\text{R}^3)_2$, NO_2 , CN , COOR^3 , $\text{CON}(\text{R}^3)_2$, NR^3COA , $\text{NR}^3\text{CON}(\text{R}^3)_2$, $\text{NR}^3\text{SO}_2\text{A}$, COR^3 , $\text{SO}_2\text{N}(\text{R}^3)_2$, $\text{S}(\text{O})_n\text{A}$, $-\text{C}(\text{R}^4)_2]_n\text{-COOR}^3$ or $-\text{O}[\text{C}(\text{R}^4)_2]_o\text{-COOR}^3$,
- Ar' is phenyl, naphthyl or biphenyl, each of which is unsubstituted or monosubstituted, disubstituted or trisubstituted by Hal, A, OR^4 , $\text{N}(\text{R}^4)_2$, NO_2 , CN , COOR^4 , $\text{CON}(\text{R}^4)_2$, NR^4COA , $\text{NR}^4\text{CON}(\text{R}^4)_2$, $\text{NR}^4\text{SO}_2\text{A}$, COR^4 , $\text{SO}_2\text{N}(\text{R}^4)_2$, $\text{S}(\text{O})_n\text{A}$, $-\text{C}(\text{R}^4)_2]_n\text{-COOR}^4$ or $-\text{O}[\text{C}(\text{R}^4)_2]_o\text{-COOR}^4$,
- Het is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic ring having from 1 to 4 N, O and/or S atoms which may be unsubstituted or monosubstituted,

disubstituted or trisubstituted by Hal, A, $-\text{C}(\text{R}^4)_2\text{-Ar}$,
 $-\text{C}(\text{R}^4)_2\text{-Het}'$, $-\text{C}(\text{R}^4)_2\text{-cycloalkyl}$, OR^3 , $\text{N}(\text{R}^3)_2$,
 $\text{NR}^3\text{CON}(\text{R}^3)_2$, NO_2 , CN , $-\text{C}(\text{R}^4)_2\text{-COOR}^3$,
 $-\text{C}(\text{R}^4)_2\text{-CON}(\text{R}^3)_2$, NR^3COA , $\text{NR}^3\text{SO}_2\text{A}$, COR^3 , SO_2NR^3 ,
 $\text{S}(\text{O})_m\text{A}$ and/or carbonyl oxygen,

Het' is a monocyclic or bicyclic, saturated, unsaturated or aromatic heterocyclic ring having from 1 to 4 N, O and/or S atoms which may be unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, $=\text{S}$, $=\text{N}(\text{R}^4)_2$, Hal, A, OR^4 , $\text{N}(\text{R}^4)_2$, NO_2 , CN , COOR^4 , $\text{CON}(\text{R}^4)_2$, NR^4COA , $\text{NR}^4\text{CON}(\text{R}^4)_2$, $\text{NR}^4\text{SO}_2\text{A}$, COR^4 , SO_2NR^4 and/or $\text{S}(\text{O})_n\text{A}$,

Hal is F, Cl, Br or I,

n is 0, 1 or 2,

o is 1, 2 or 3,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

2. Compounds according to Claim 1, in which

D is a monocyclic or bicyclic, aromatic carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or disubstituted by Hal,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

3. Compounds according to Claim 1 or 2, in which

D is phenyl, pyridyl, thienyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

4. Compounds according to one or more of Claims 1-3, in which

R^1, R^2 are each, independently of one another, H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy, NHCOA, NHSO₂A, OCH₂COOA or OCH₂COOH,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

5. Compounds according to one or more of Claims 1-4, in which

G is (CH₂)_n, (CH₂)_nNH-, -CH=CH- or -CH=CH-CH=CH-, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

6. Compounds according to one or more of Claims 1-5, in which

X is -[C(R⁴)₂]_nCONR³[C(R⁴)₂]_n-, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

7. Compounds according to one or more of Claims 1-6, in which

X is -CONH- or -CON(CH₂COOA)-, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

8. Compounds according to one or more of Claims 1-7, in which

Y is cycloalkylene, Het-diyl or Ar-diyl, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

9. Compounds according to Claims 1-8,

in which

Y is pyridinediyl, piperidinediyl, cyclohexylene, or phenylene which is unsubstituted or monosubstituted or disubstituted by A, OA, Cl, F, COOCH₃, COOH, phenoxy or aminocarbonyl, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

10. Compounds according to one or more of Claims 1-9,

in which

T is a monocyclic, saturated or unsaturated heterocyclic ring having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by =O, =S or =NH and may be monosubstituted or disubstituted by Hal, A and/or OA, and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

11. Compounds according to one or more of Claims 1-10,

in which

T is piperidin-1-yl, pyrrolidin-1-yl, pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl, imidazolidinyl, thiazolyl or 1,4-oxazepanyl, each of which is monosubstituted or disubstituted by =O or =NH and where the radicals may also be monosubstituted or disubstituted by Hal, A and/or OA,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

12. Compounds according to one or more of Claims 1-11,

in which

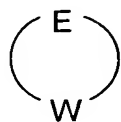
Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OA, SO₂A, COOR², SO₂NH₂, CN, COOA, COOH or phenoxy,
and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

13. Compounds according to one or more of Claims 1-12,
in which

- 10 D is a monocyclic or bicyclic, aromatic carbocyclic or heterocyclic ring having from 0 to 4 N, O and/or S atoms which is unsubstituted or monosubstituted or disubstituted by Hal,
- 15 R¹, R² are each, independently of one another, H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy, NHCOA, NHSO₂A, OCH₂COOA or OCH₂COOH,
- 20 R¹ and R² together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
- R³ is H, A, phenyl, benzyl or [C(R⁴)₂]_nCOOA,
- R⁴ is H or A,
- W is N, CR³ or an sp²-hybridised carbon atom,
- 25 E together with W is a 3- to 7-membered saturated carbocyclic or heterocyclic ring having from 0 to 3 N, from 0 to 2 O and/or from 0 to 2 S atoms, which may contain a double bond,
- 30 G is (CH₂)_n, (CH₂)_nNH-, -CH=CH- or -CH=CH-CH=CH-,
- X is -[C(R⁴)₂]_nCONR³[C(R⁴)₂]_n-,
- Y is cycloalkylene, Het-diyl or Ar-diyl,
- Ar is phenyl which is unsubstituted or monosubstituted or disubstituted by Hal, A, OA, SO₂A, COOR², SO₂NH₂, CN, COOA, COOH or phenoxy,
- 35

- 5 T is a monocyclic, saturated or unsaturated heterocyclic ring having 1 to 2 N and/or O atoms which is monosubstituted or disubstituted by =O, =S or =NH and may be monosubstituted or disubstituted by Hal, A and/or OA,
- A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- Hal is F, Cl, Br or I,
- n is 0, 1 or 2,
- 10 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
14. Compounds according to one or more of Claims 1-13,
- 15 in which
- D is phenyl, pyridyl, thienyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,
- R¹, R² are each, independently of one another, H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy, NHCOA, NHSO₂A, OCH₂COOA or OCH₂COOH,
- 20 R¹ and R² together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
- 25 R³ is H, A or CH₂COOA,
- R⁴ is H or A,
- W is N, CR³ or an sp²-hybridised carbon atom,
- E together with W is a 3- to 7-membered saturated carbocyclic or heterocyclic ring having from 0 to 3 N, from 0 to 2 O and/or from 0 to 2 S atoms,
- 30 which may contain a double bond,
- G is (CH₂)_n, (CH₂)_nNH-, -CH=CH- or -CH=CH-CH=CH-,
- X is -CONH- or -CON(CH₂COOA)-,
- 35 Y is pyridinediyl, piperidinediyl, cyclohexylene, or phenylene which is unsubstituted or monosubstituted or

- disubstituted by A, OA, Cl, F, COOCH₃, COOH,
phenoxy or aminocarbonyl,
T is piperidin-1-yl, pyrrolidin-1-yl, pyridin-1-yl, morpholin-4-
yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, pyridazin-2-yl,
5 pyrazin-1-yl, azepan-1-yl, 2-azabicyclo[2.2.2]octan-2-yl,
imidazolidinyl, thiazolyl or 1,4-oxazepanyl, each of which
is monosubstituted or disubstituted by =O or =NH and
where the radicals may also be monosubstituted or
10 disubstituted by Hal, A and/or OA,
A is unbranched or branched alkyl having 1-10 carbon
atoms and in which 1-7 H atoms may be replaced by F,
Hal is F, Cl, Br or I,
15 n is 0, 1 or 2,
and pharmaceutically usable derivatives, solvates, salts and stereo-
isomers thereof, including mixtures thereof in all ratios.
15. Compounds according to one or more of Claims 1-14,
20 in which
D is phenyl, pyridyl or thienyl, each of which is
monosubstituted or disubstituted by Hal,
R¹ is H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4,
25 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy,
-OCOR³, NHCOA or NHSO₂A,
R² is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 car-
bon atoms,
30 R¹ and R² together are alternatively a spirocyclically bonded 3- to
6-membered carbocyclic ring,
R³ is H or A,
R⁴ is H or A,



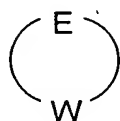
- is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,
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- G is $(\text{CH}_2)_n$ or $(\text{CH}_2)_n\text{NH}-$,
- 10 X is CONH,
- Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,
- 15 T is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo-[2.2.2]octan-2-yl, each of which is monosubstituted or disubstituted by carbonyl oxygen,
- 20 A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- Hal is F, Cl, Br or I,
- n is 0, 1 or 2;
- 25 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
16. Compounds according to one or more of Claims 1-15, in which
- 30 D is phenyl, pyridyl or thienyl, each of which is monosubstituted or disubstituted by Hal,
- R¹ is H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy,
- 35 -OCOR³, NHCOA or NHSO₂A,

R^2 is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

R^1 and R^2 together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,

R^3 is H or A,

R^4 is H or A,



is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-

3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-

pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-

3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or

azetidine-1,2-diyl,

G is $(CH_2)_n$ or $(CH_2)_nNH-$,

X is CONH,

Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,

A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F;

Hal is F, Cl, Br or I,

n is 0, 1 or 2;

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

17. Compounds according to one or more of Claims 1-16, in which

X is $-[C(R^4)_2]_nCONR^3[C(R^4)_2]_n-$ or $-[C(R^4)_2]_nCO[C(R^4)_2]_n-$,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

18. Compounds according to one or more of Claims 1-17,
in which

X is CONH or COCH₂,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

19. Compounds according to one or more of Claims 1-18,
in which

D is phenyl, pyridyl or thienyl, each of which is monosubstituted or disubstituted by Hal,

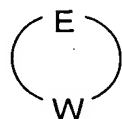
R¹ is H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy, -OCOR³, NHCOA or NHSO₂A,

R² is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

R¹ and R² together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,

R³ is H or A,

R⁴ is H or A,



is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-

3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1H-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,

G is (CH₂)_n or (CH₂)_nNH₂,

X is CONH or COCH₂,

Y is 1,3- or 1,4-phenylene which is unsubstituted or monosubstituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

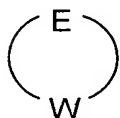
T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,
 A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
 5 Hal is F, Cl, Br or I,
 n is 0, 1 or 2,
 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

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20. Compounds according to one or more of Claims 1-19, in which

D is phenyl, pyridyl or thienyl, each of which is monosubstituted or disubstituted by Hal,
 15 R¹ is H, =O, COOR³, OH, OA, NH₂, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N₃, ethynyl, vinyl, allyloxy, -OCOR³, NHCOA, NHSO₂A, H-C≡C-CH₂-, CH₃-C≡C-CH₂-O-, -O-CH₂-CH(OH)-CH₂OH,
 20 -O-CH₂-CH(OH)-CH₂NH₂ or -O-CH₂-CH(OH)-CH₂Het',
 R² is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,
 R¹ and R² together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
 25 R³ is H or A,
 R⁴ is H or A,

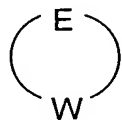
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is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1H-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,
 G is (CH₂)_n or (CH₂)_nNH-,

- X is CONH or COCH₂,
- Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,
- 5 T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,
- Het' is a saturated 3-6-membered heterocyclic ring having from 1 to 3 N and/or O atoms, which may be unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, Hal, A, OH, NH₂, NO₂, CN, COOA or CONH₂,
- 10 A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- 15 Hal is F, Cl, Br or I,
- n is 0, 1 or 2,
- and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.
- 20 21. Compounds according to one or more of Claims 1-20, in which
- D is phenyl, pyridyl or thienyl, each of which is monosubstituted or disubstituted by Hal,
- 25 R¹ is ethynyl, vinyl, allyloxy, CH₃-C≡C-CH₂-O-, -O-CH₂-CH(OH)-CH₂OH, -O-CH₂-CH(OH)-CH₂NH₂ or -O-CH₂-CH(OH)-CH₂Het',
- R² is H or OH,
- 30 R¹ and R² together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
- R³ is H or A,
- R⁴ is H or A,



is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-

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3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1*H*-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,

G

is (CH₂)_n or (CH₂)_nNH-,

10

X

is CONH, CO, COO or COCH₂,

Y

is 1,3- or 1,4-phenylene which is unsubstituted or monosubstituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

15

T

is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo-[2.2.2]octan-2-yl, each of which is monosubstituted or disubstituted by carbonyl oxygen or OA,

20

Het'

is a saturated 3-6-membered heterocyclic ring having from 1 to 3 N and/or O atoms, which may be unsubstituted or monosubstituted or disubstituted by carbonyl oxygen, Hal, A, OH, NH₂, NO₂, CN, COOA or CONH₂,

25

A

is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,

Hal

is F, Cl, Br or I,

n

is 0, 1 or 2,

30

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

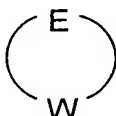
22. Compounds according to one or more of Claims 1-21,

in which

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D

is phenyl, pyridyl, thienyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,

- 5
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- R^1 is H, =O, COOR^3 , OH, OA, NH_2 , alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N_3 , ethynyl, vinyl, allyloxy, NHCOA , NHSO_2A , OCH_2COOA or OCH_2COOH ,
- R^2 is H, =O, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,
- R^1 and R^2 together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,
- R^3 is H or A,
- R^4 is H or A,
-  is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl, thiazolidine-3,4-diyl, 2,5-dihydro-1H-pyrrole-1,5-diyl, 1,3-dioxolane-4,5-diyl, 1,3-oxazinane-3,4-diyl, piperazine-1,4-diyl, tetrahydrofuran-3,4-diyl or azetidine-1,2-diyl,
- G is $(\text{CH}_2)_n$, $(\text{CH}_2)_n\text{NH}-$, $-\text{CH}=\text{CH}-$ or $-\text{CH}=\text{CH}-\text{CH}=\text{CH}-$,
- X is CONH , COCH_2 or $-\text{CON}(\text{CH}_2\text{COOA})-$,
- Y is pyridinediyl, piperidinediyl, cyclohexylene, or phenylene which is unsubstituted or monosubstituted or disubstituted by A, OA, Cl, F, COOCH_3 , COOH , phenoxy or aminocarbonyl,
- T is morpholin-4-yl which is monosubstituted or disubstituted by carbonyl oxygen,
- A is unbranched or branched alkyl having 1-10 carbon atoms and in which 1-7 H atoms may be replaced by F,
- Hal is F, Cl, Br or I,
- n is 0, 1 or 2,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

23. Compounds according to Claim 1, selected from the group consisting of

- 5 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
10 1-N-[(4-chlorophenyl)]-2-N-[[3-fluoro-4-(3-oxomorpholin-4-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
15 1-N-[(4-chlorophenyl)]-2-N-[[3-trifluoromethyl-4-(3-oxomorpholin-4-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]]-(R)-piperidine-1,2-dicarboxamide,
1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2H-pyridin-1-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
20 1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2H-pyrazin-1-yl)phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,
1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(R)-2,5-dihydropyrrole-1,2-dicarboxamide,
25 N-[4-(3-oxomorpholin-4-yl)phenyl]-(R)-1-(5-chlorothiophene-2-carbonyl)pyrrolidine-2-carboxamide,
N-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]-(R)-1-(5-chlorothiophene-2-carbonyl)pyrrolidine-2-carboxamide,
30 3-N-[(4-chlorophenyl)]-4-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(R)-oxazolidine-3,4-dicarboxamide,
3-N-[(4-chlorophenyl)]-4-N-[[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]]-(R)-oxazolidine-3,4-dicarboxamide,
35 3-N-[(4-chlorophenyl)]-4-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[4-(2-oxo-2H-pyridin-1-yl)phenyl]]-(R)-oxazolidine-3,4-dicarboxamide,

5 3-N-[(4-chlorophenyl)-4-N-[[4-(2-oxo-2H-pyridin-1-yl)phenyl]]-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[3-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

10 3-N-[(4-chlorophenyl)-4-N-[[3-chloro-4-(3-oxomorpholin-4-yl)-phenyl]]-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(4R,5R)-5-methyloxazolidine-3,4-dicarboxamide,

15 3-N-[(4-chlorophenyl)-4-N-[[4-(2-oxo-2H-pyrazin-1-yl)phenyl]]-(4R,5S)-5-methyloxazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[4-(2-oxo-2H-pyrazin-1-yl)phenyl]]-(R)-oxazolidine-3,4-dicarboxamide,

20 3-N-[(4-chlorophenyl)-4-N-[[3-chloro-4-(2-oxo-2H-pyridin-1-yl)phenyl]]-(R)-oxazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(S)-thiazolidine-3,4-dicarboxamide,

25 3-N-[(4-chlorophenyl)-4-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(S)-1,1-dioxo-1 λ^6 -thiazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(S)-thiazolidine-3,4-dicarboxamide,

30 3-N-[(4-chlorophenyl)-4-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(S)-1,1-dioxo-1 λ^6 -thiazolidine-3,4-dicarboxamide,

3-N-[(4-chlorophenyl)-4-N-[[4-(2-oxo-2H-pyridin-1-yl)phenyl]]-(R)-thiazolidine-3,4-dicarboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxamide,

35 N-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]-3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxamide,

N-[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]-3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxamide,

1-N-[(5-chloropyridin-2-yl)]-2-N-[[4-(2-oxo-2*H*-pyridin-1-yl)-phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

5

1-N-[(5-chloropyridin-2-yl)]-2-N-[[4-(3-oxomorpholin-4-yl)-phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(5-chloropyridin-2-yl)]-2-N-[[4-(2-oxo-2*H*-pyrazin-1-yl)-phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

10

1-N-[(5-chloropyridin-2-yl)]-2-N-[[3-fluoro-4-(2-oxo-2*H*-pyridin-1-yl)phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(5-chloropyridin-2-yl)]-2-N-[[4-(2-oxo-2*H*-pyridin-1-yl)-phenyl]]-(*R*)-4,4-dimethoxypyrrolidine-1,2-dicarboxamide,

15

1-N-[(5-chloropyridin-2-yl)]-2-N-[[4-(3-oxomorpholin-4-yl)-phenyl]]-(*R*)-4,4-dimethoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]]-(*R*)-4,4-dimethoxypyrrolidine-1,2-dicarboxamide,

20

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

25

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

30

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxopyrazin-1-yl)phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-fluoro-4-(2-oxo-2*H*-pyridin-1-yl)-phenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

35

1-N-[(4-chlorophenyl)]-2-N-[[3-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(2*R*,3*R*)-3-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(2R,3S)-3-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4S)-4-hydroxypyrrolidine-1,2-dicarboxamide,

5 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2S,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
3,4-dihydroxypyrrolidine-1,2-dicarboxamide,

10 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4S)-4-azidopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4S)-4-aminopyrrolidine-1,2-dicarboxamide,

15 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-azidopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-aminopyrrolidine-1,2-dicarboxamide,

20 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4S)-4-acetaminopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-acetaminopyrrolidine-1,2-dicarboxamide,

25 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4S)-4-methylsulfonylaminopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-methylsulfonylaminopyrrolidine-1,2-dicarboxamide,

30 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,

35 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-propoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-allyloxypyrrolidine-1,2-dicarboxamide,

(3R,5R)-1-(4-chlorophenylcarbamoyl)-5-[4-(3-oxomorpholin-4-yl)phenylcarbamoyl]pyrrolidin-3-yl isobutyrate,

(3R,5R)-1-(4-chlorophenylcarbamoyl)-5-[4-(3-oxomorpholin-4-yl)phenylcarbamoyl]pyrrolidin-3-yl propionate,

5 (3R,5R)-1-(4-chlorophenylcarbamoyl)-5-[4-(3-oxomorpholin-4-yl)phenylcarbamoyl]pyrrolidin-3-yl acetate,

4-N-[(4-chlorophenyl)]-5-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-1,3-dioxolane-4,5-dicarboxamide,

10 4-N-[(4-chlorophenyl)]-5-N-[[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]]-1,3-dioxolane-4,5-dicarboxamide,

4-N-[(4-chlorophenyl)]-5-N-[[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]]-1,3-dioxolane-4,5-dicarboxamide,

15 4-N-[(4-chlorophenyl)]-5-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-1,3-dioxolane-2,2-dimethyl-4,5-dicarboxamide,

4-N-[(4-chlorophenyl)]-5-N-[[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]]-1,3-dioxolane-2,2-dimethyl-4,5-dicarboxamide,

20 4-N-[(4-chlorophenyl)]-5-N-[[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]]-1,3-dioxolane-2,2-dimethyl-4,5-dicarboxamide,

1-N-[4-chlorophenyl]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-1-BOC-piperazine-1,2-dicarboxamide,

25 1-N-[4-chlorophenyl]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-piperazine-1,2-dicarboxamide,

1-N-[4-chlorophenyl]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-1,3-oxazinane-3,4-dicarboxamide,

30 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4S)-4-ethynyl-4-hydroxypyrrolidine-1,2-dicarboxamide,

6-N-[(4-chlorophenyl)]-7-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-4-oxa-6-azaspiro[2.4]heptane-6,7-dicarboxamide,

35 1-N-[(6-chloropyridin-3-yl)]-2-N-[[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(6-chloropyridin-3-yl)]-2-N-[[4-(2-oxo-2*H*-pyrazin-1-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4S)-4-acetaminopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4S)-4-butylsulfonylaminopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(R)-4-oxopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4S)-4-aminopyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(S)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[2-(4-chlorophenyl)acetyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-(4-chlorobenzoyl)-4-hydroxypyrrolidine-2-carboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2H-pyridin-1-yl)phenyl]]-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2H-pyrazin-1-yl)phenyl]]-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4S)-4-(2-methylpropanoylamino)pyrrolidine-1,2-dicarboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-(1-1H-indol-3-yl-methanoyl)-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-(1-1H-indol-6-yl-methanoyl)-4-hydroxypyrrolidine-2-carboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]]-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]]-(2R,4*S*)-4-ethynyl-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2*H*-pyrazin-1-yl)phenyl]]-(2R,4*S*)-4-ethynyl-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2*H*-pyridin-1-yl)phenyl]]-4,4-difluoro-(*R*)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(2-oxo-2*H*-pyridin-1-yl)-phenyl]]-(2R,4*R*)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(2-oxo-2*H*-pyridin-1-yl)-phenyl]]-(*R*)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(2-oxo-2*H*-pyridin-1-yl)-phenyl]]-(2R,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

2-N-[(4-chlorophenyl)]-1-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(*R*)-pyrrolidine-1,2-dicarboxamide,

2-N-[(4-chlorophenyl)]-1-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(*S*)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-3-methoxy-2*H*-pyridin-1-yl)phenyl]]-(2R,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-3-methoxy-2*H*-pyridin-1-yl)phenyl]]-(*R*)-pyrrolidine-1,2-dicarboxamide,

N-(4-chlorophenyl)-(R)-1-{2-[4-(3-oxomorpholin-4-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(S)-1-{2-[4-(3-oxomorpholin-4-yl)phenyl]-acetyl}pyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(2R,4R)-1-{2-[4-(3-oxomorpholin-4-yl)-phenyl]acetyl}-4-methoxypyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(2R,4S)-1-{2-[4-(3-oxomorpholin-4-yl)-phenyl]acetyl}-4-methoxypyrrolidine-2-carboxamide,

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N-(4-chlorophenyl)-(2S,4R)-1-{2-[4-(3-oxomorpholin-4-yl)-phenyl]acetyl}-4-methoxypyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(S)-1-{2-[4-(2-oxo-1H-pyridin-1-yl)phenyl]acetyl}pyrrolidine-2-carboxamide,

10

N-(4-chlorophenyl)-(S)-1-{2-[4-(2-oxopyrrolidin-1-yl)phenyl]acetyl}pyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(R)-1-{2-[4-(2-oxopyrrolidin-1-yl)phenyl]acetyl}pyrrolidine-2-carboxamide,

15

N-(4-chlorophenyl)-(R)-1-[4-(2-oxopiperidin-1-yl)benzoyl]pyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(R)-1-[4-(2-oxopiperidin-1-yl)phenoxy]carbonylpyrrolidine-2-carboxamide,

20

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-2H-pyrazin-1-yl)phenyl]]-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-fluoro-4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-ethoxypyrrolidine-1,2-dicarboxamide,

25

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-(prop-2-ynyloxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-(but-2-ynyloxy)pyrrolidine-1,2-dicarboxamide,

30

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-(2,3-dihydroxypropoxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-(2-hydroxy-3-pyrrolidin-1-ylpropoxy)pyrrolidine-1,2-dicarboxamide,

35

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-(2-oxooxazolidin-5-ylmethoxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-(3-amino-2-hydroxypropoxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-1*H*-pyrazin-1-yl)phenyl]]-
(R)-2,5-dihydropyrrole-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-oxo-1*H*-pyridin-1-yl)phenyl]]-
(R)-2,5-dihydropyrrole-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[3-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(R)-2,5-dihydropyrrole-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(R)-2,5-dihydropyrrole-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2S,3S)-3-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2S,4S)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-methoxycarbonyl-4-(3-oxomor-
pholin-4-yl)phenyl]]-(2R,4R)-3-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-carboxy-4-(3-oxomorpholin-4-yl)-
phenyl]]-(2R,4R)-3-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,3S,4R)-3,4-dihydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(2R,4R)-4-allyloxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(2R,4R)-4-(prop-2-ynyloxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(2R,4S)-4-(prop-2-ynyloxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-(methoxycarbonylmethoxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-
(2R,4R)-4-(carboxymethoxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-bromophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-
phenyl]]-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-(3-oxomorpholin-4-yl)-phenyl]]-(2R,4R)-4-(2,3-dihydroxypropoxy)pyrrolidine-1,2-dicarboxamide,

5 1-N-[(4-chlorophenyl)]-2-N-{N-methoxycarbonylmethyl-N'-[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-methoxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)cyclohexan-1-yl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

10 1-N-[(4-chlorophenyl)]-2-N-[[4-(2-iminopyrrolidin-1-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 442;

1-N-[(4-chlorophenyl)]-2-N-[[3-methyl-4-(2-iminopyrrolidin-1-yl)-phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 456;

15 1-N-[(4-chlorophenyl)]-2-N-[4-{2-[(E)-cyanimino]imidazolidin-1-yl}phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 468;

1-N-[(4-chlorophenyl)]-2-N-[[4-(2-imino-5-methylthiazol-3-yl)-phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 473;

20 1-N-[(4-chlorophenyl)]-2-N-[[2-aminocarbonyl-4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide, ESI 502;

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxy-2-methylpyrrolidine-1,2-dicarboxamide,

25 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-thiophen-3-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

30 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(2E,4E)-5-phenylpenta-2,4-dienyloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-methylfuran-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

35 N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-thiophen-2-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1H-imidazol-4-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

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N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-methoxypyrrolidine-2-carboxamide,

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N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-chlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(3,4-dichlorophenyl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

15

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorofuran-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-chlorothiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

20

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

25

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,

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N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,

35

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[2-fluoro-4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

5

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-3-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-4-ylacryloyl]-4-hydroxypyrrolidine-2-carboxamide,

10

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-pyridin-4-ylacryloyl]-4-ethoxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-1*H*-imidazol-4-ylacryloyl]-4-methoxypyrrolidine-2-carboxamide,

15

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-bromothiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(4-bromothiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

20

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-bromothiophen-2-yl)acryloyl]-4-hydroxypyrrolidine-2-carboxamide,

N-[4-(3-oxomorpholin-4-yl)phenyl]-(2R,4R)-1-[(E)-3-(5-bromothiophen-2-yl)acryloyl]-4-ethoxypyrrolidine-2-carboxamide,

25

N-(4-chlorophenyl)-(R)-1-[4-(2-oxopiperidin-1-yl)benzoyl]pyrrolidine-2-carboxamide,

N-(4-chlorophenyl)-(S)-1-[4-(2-oxopiperidin-1-yl)benzoyl]pyrrolidine-2-carboxamide,

30

1-N-[(4-chlorophenyl)]-2-N-[[4-(5-oxo-1,4-oxazepan-4-yl)-phenyl]]-(R)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(5-oxo-1,4-oxazepan-4-yl)-phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

35

1-N-[(4-chlorophenyl)]-2-N-[[4-((S)-2-methyl-3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-((S)-2-methyl-3-oxomorpholin-4-yl)phenyl]]-(2R)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-((R)-2-methyl-3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-((R)-2-methyl-3-oxomorpholin-4-yl)phenyl]]-(2R)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)-2-phenoxyphenyl]]-(2R)-pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-fluoro-4-((R)-2-methyl-3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-3-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-piperidine-1,3-dicarboxamide,

1-N-[(4-chlorophenyl)]-3-N-[[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]]piperidine-1,3-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-(2-methoxyethoxy)pyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxo-1,4-oxazepan-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-methyl-4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

24. Pyrrolidinecarboxylic acid derivatives selected from the group consisting of

1-N-[(4-chlorophenyl)]-2-N-[(1'-methyl-[1,4']bipiperidiny-4-yl)]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[(3,4,5,6-tetrahydro-2*H*-1,4'-bipyridinyl-4-yl)]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[(3,4,5,6-tetrahydro-2*H*-1,4'-bipyridinyl-4-yl)]-(2*R*,4*R*)-4-ethoxypyrrolidine-1,2-dicarboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-4-hydroxy-2-(4-pyridin-4-yl)piperazine-1-carbonylpyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-4-hydroxy-2-[4-(2-methoxyphenyl)piperazine-1-carbonyl]pyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-2-[4-(4-fluorophenyl)piperazine-1-carbonyl]-4-hydroxypyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-4-hydroxy-2-[4-hydroxy-4-(4-methoxyphenyl)piperidine-1-carbonyl]pyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-4-hydroxy-2-(4-pyridin-2-yl)piperazine-1-carbonylpyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-2-[4-(4-ethylpiperazin-1-yl)piperidine-1-carbonyl]-4-hydroxypyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-2-[4-(4,6-dimethylpyrimidin-2-yl)piperazine-1-carbonyl]-4-hydroxypyrrolidine-1-carboxamide,

N-(4-chlorophenyl)-(2*R*,4*R*)-4-hydroxy-2-[4-(1-methylpiperidin-4-yl)piperazine-1-carbonyl]pyrrolidine-1-carboxamide,

1-N-[(4-chlorophenyl)]-2-N-[[2-(2-dimethylaminoethoxy)-4-morpholin-4-ylphenyl]]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[(2-ethoxy-4-morpholin-4-ylphenyl)]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

1-N-[(4-chlorophenyl)]-2-N-[(4-morpholin-4-yl-2-propoxyphenyl)]-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

25. Cyclopentanecarboxylic acid derivatives selected from the group consisting of

N-[4-(3-oxomorpholin-4-yl)phenyl]-(rac)-2-[3-(4-chlorophenyl)-ureido]cyclopentanecarboxamide,

N-[3-methyl-4-(3-oxomorpholin-4-yl)phenyl]-(rac)-2-[3-(4-chlorophenyl)ureido]cyclopentanecarboxamide,

and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios.

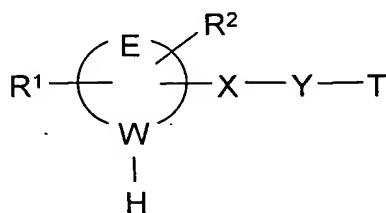
26. Process for the preparation of compounds of the formula I according to Claims 1-23 and pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, characterised in that

a) for the preparation of compounds of the formula I in which

W is N and

G is NH,

a compound of the formula II

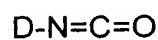


II

in which

R¹, R², E, X, Y and T are as defined in Claim 1,
and W is N,

is reacted with a compound of the formula III



III

in which

D is as defined in Claim 1,

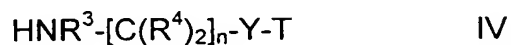
or

5

b) for the preparation of compounds of the formula I in which
X is $-[C(R^4)_2]_nCONR^3[C(R^4)_2]_n-$,

a compound of the formula IV

10

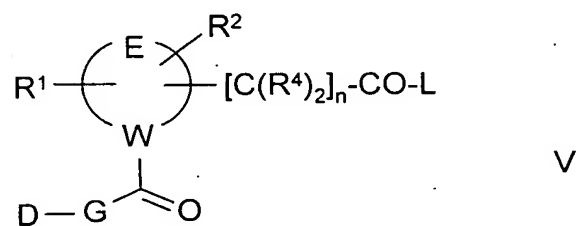


in which R^3 , n, Y and T are as defined in Claim 1,

15

is reacted with a compound of the formula V

20



in which

25

L is Cl, Br, I or a free or reactively functionally modified OH group,
and

R^1 , R^2 , R^4 , D, E, G, W and n are as defined in Claim 1,

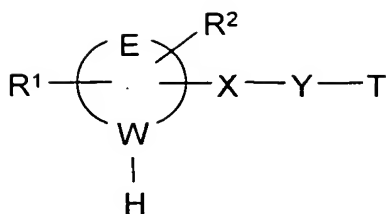
30

or

c) for the preparation of compounds of the formula I in which W is
N,

35

a compound of the formula II



II

in which

R^1 , R^2 , E, X, Y and T are as defined in Claim 1,
and W is N,

is reacted with a compound of the formula VI



VI

in which D and G are as defined in Claim 1, and

L is Cl, Br, I or a free or reactively functionally modified OH group,

and/or

a base or acid of the formula I is converted into one of its salts.

27. Compounds of the formula I according to one or more of Claims 1 to 23 and the compounds of Claims 24 and 25 as inhibitors of coagulation factor Xa.

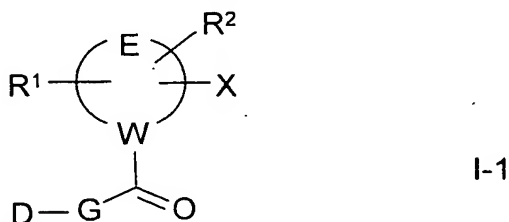
28. Compounds of the formula I according to one or more of Claims 1 to 23 and the compounds of Claims 24 and 25 as inhibitors of coagulation factor VIIa.

29. Medicament comprising at least one compound of the formula I according to one or more of Claims 1 to 23 or a compound of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, and, if desired, excipients and/or adjuvants.

- 5 30. Medicament comprising at least one compound of the formula I according to one or more of Claims 1 to 23 or a compound of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates and stereoisomers thereof, including mixtures thereof in all ratios, and at least one further medicament active ingredient.
- 10 31. Use of compounds according to one or more of Claims 1 to 23 or the compounds of Claims 24 and 25 and/or physiologically acceptable salts, salts and solvates thereof for the preparation of a medicament for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases.
- 15 32. Set (kit) consisting of separate packs of
- 20 (a) an effective amount of a compound of the formula I according to one or more of Claims 1 to 23 or a compound of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, and
- 25 (b) an effective amount of a further medicament active ingredient.
- 30 33. Use of compounds of the formula I according to one or more of Claims 1 to 23 or of compounds of Claims 24 and 25 and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, for the preparation of a medicament for the treatment of thromboses, myocardial infarction, arteriosclerosis, inflammation, apoplexy, angina pectoris, restenosis after angioplasty, claudicatio intermittens, migraine, tumours, tumour diseases and/or tumour metastases,
- 35

in combination with at least one further medicament active ingredient.

34. Intermediate compounds of the formula I-1

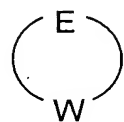


10 in which

D is phenyl, -pyridyl, thienyl, furyl or imidazolyl, each of which is monosubstituted or disubstituted by Hal,

R¹ is H, OH, OA, alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms or ethynyl,

15 R² is H, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,



is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4-

or 3,5-diyl,

G is (CH₂)_n, (CH₂)_nNH-, -CH=CH- or -CH=CH-CH=CH-,

X is COOH,

25 A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

Hal is F, Cl, Br or I,

n is 0, 1 or 2,

and isomers and salts thereof.

30 35. Compounds according to Claim 34, selected from the group consisting of

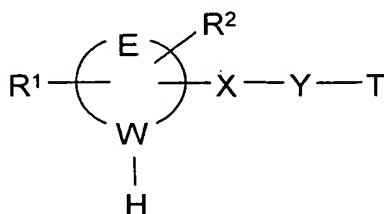
3-(4-chlorophenylcarbamoyl)oxazolidine-4-carboxylic acid,

3-(5-chlorothiophene-2-carbonyl)oxazolidine-5-carboxylic acid,

35 and isomers and salts thereof.

36. Intermediate compounds selected from the group consisting of
 (2R,4S)-BOC-4-ethynyl-4-hydroxy-pyrrolidine-2-carboxylic acid,
 (2R,4R)-BOC-4-ethynyl-4-hydroxy-pyrrolidine-2-carboxylic acid,
 alkyl (2R,4S)-BOC-4-ethynyl-4-hydroxypyrrolidine-2-carboxylate,
 alkyl (2R,4R)-BOC-4-ethynyl-4-hydroxypyrrolidine-2-carboxylate,
 where alkyl has 1, 2, 3, 4, 5 or 6 carbon atoms,
 and isomers and salts thereof.

37. Intermediate compounds of the formula I-2



I-2

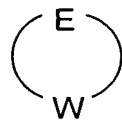
in which

R^1 is H, =O, COOR^3 , OH, OA, NH_2 , alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N_3 , ethynyl, vinyl, allyloxy, NHCOA , $\text{NH}\text{SO}_2\text{A}$, OCH_2COOA or OCH_2COOH ,

R^2 is H, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

R^1 and R^2 together are alternatively a spirocyclically bonded 3- to 6-membered carbocyclic ring,

R^3 is H or A,



is pyrrolidine-1,2-diyl, piperidine-1,2-diyl, oxazolidine-3,4- or 3,5-diyl,

X is CONH,

Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

T is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo[2.2.2]octan-2-yl, each of which is mono-substituted or disubstituted by carbonyl oxygen,

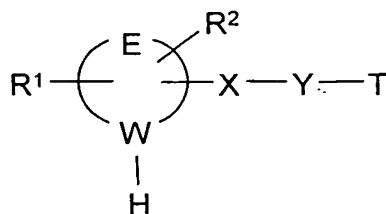
A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

Hal is F, Cl, Br or I,

n is 0, 1 or 2,

and isomers and salts thereof.

38. Compounds according to Claim 37 of the formula I-2a



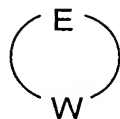
I-2a

in which

R^1 is H, =O, COOR^3 , OH, OA, NH_2 , alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms, N_3 , ethynyl, vinyl, allyloxy, NHCOA , NHSO_2A , OCH_2COOA or OCH_2COOH ,

R^2 is H, OH, OA or alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,

R^3 is H or A,



is pyrrolidine-1,2-diyl,

X is CONH,

Y is 1,3- or 1,4-phenylene which is unsubstituted or mono-substituted or disubstituted by methyl, trifluoromethyl, ethyl, propyl, Cl or F,

- 5 T is piperidin-1-yl, pyrrolidin-1-yl, 1*H*-pyridin-1-yl, morpholin-4-yl, piperazin-1-yl, 1,3-oxazolidin-3-yl, 2*H*-pyridazin-2-yl, pyrazin-1-yl, azepan-1-yl or 2-azabicyclo[2.2.2]octan-2-yl, each of which is monosubstituted or disubstituted by carbonyl oxygen,
- A is alkyl having 1, 2, 3, 4, 5 or 6 carbon atoms,
- Hal is F, Cl, Br or I,
- n is 0, 1 or 2,
- 10 and isomers and salts thereof.
39. Compounds according to Claim 38, selected from the group consisting of
- 15 *N*-[4-(3-oxomorpholin-4-yl)phenyl]-(*S*)-pyrrolidine-2-carboxamide,
- N*-[4-(3-oxomorpholin-4-yl)phenyl]-(*R*)-pyrrolidine-2-carboxamide,
- N*-[4-(3-oxomorpholin-4-yl)phenyl]-(2*R*,4*R*)-4-hydroxypyrrolidine-2-carboxamide,
- 20 *N*-[4-(3-oxomorpholin-4-yl)phenyl]-4-hydroxypyrrolidine-2-carboxamide,
- N*-[4-(3-oxomorpholin-4-yl)phenyl]-(*R*)-4,4-dimethoxypyrrolidine-2-carboxamide,
- N*-[4-(3-oxomorpholin-4-yl)phenyl]-(2*R*,4*R*)-4-methoxypyrrolidine-2-carboxamide,
- 25 and isomers and salts thereof.
40. Medicament according to Claim 30, comprising 1-*N*-[(4-chlorophenyl)]-2-*N*-{[4-(3-oxomorpholin-4-yl)phenyl]}-(2*R*,4*R*)-4-hydroxypyrrolidine-1,2-dicarboxamide and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, and aspirin.
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41. Use according to Claim 33, comprising 1-N-[(4-chlorophenyl)]-2-N-[[4-(3-oxomorpholin-4-yl)phenyl]]-(2R,4R)-4-hydroxypyrrolidine-1,2-dicarboxamide and/or pharmaceutically usable derivatives, solvates, salts and stereoisomers thereof, including mixtures thereof in all ratios, in combination with aspirin.

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